Claims

1 A pharmaceutical composition for intramammary administration to a non-human mammal, comprising an antibacterial agent, prednisolone and a pharmaceutically acceptable carrier, characterised in that the composition comprises at least 20 mg of prednisolone / unit dose.

2 The composition according to claim 1, characterised in that it comprises the prednisolone in an amount of 20 to 40 mg / unit dose.

3 The composition according to claim 2, characterised in that it comprises the prednisolone in an amount of 20 to 30 mg / unit dose.

4 The composition according to any of claims 1 to 3, characterised in that the antibacterial agent is a cephalosporin.

5 The composition according to claim 4, characterised in that the cephalosporin is cephapirin.

6 The composition according to claim 4, characterised in that the cephalosporin is cefquinome.

7 The composition according to any of claims 1 to 6, characterised in that it comprises the antibacterial agent in an amount of 10 to 500 mg/ unit dose.

8 A process for preparing a pharmaceutical composition as claimed in any of claims 1 to 7 comprising the steps of mixing an oil and optionally additives and suspending the antibacterial agent and the prednisolone in the carrier.

9 Use of an antibacterial agent and prednisolone for the manufacture of a medicament comprising at least 20 mg of prednisolone / unit dose for the treatment of mastitis in non- human mammals.